



# Salinity-Growth Response and Ichthyotoxic Potency of the Chilean *Pseudochattonella verruculosa*

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Despite salmon farmers suffering the worst damage from a harmful algal bloom in Chile's history (US\$800M) due to a massive outbreak of the dictyochophyte *Pseudochattonella verruculosa* in 2016 ( $\sim 7000$ – $20,000$  cells  $\text{ml}^{-1}$ ), the effect of environmental drivers and the potency of lytic toxins produced by the local clones of this species remain still largely unexplored. Based on the drastic oceanographic anomalies observed in the Chilean fjords during the 2016-El Niño “Godzilla” event, the role of salinity (15 to 35 psu) on *Pseudochattonella* cell growth and cytotoxicity was studied by culturing, scanning electron microscopy (SEM) and using a rainbow trout cell line RTgill-W1 assay to define: (1) vegetative growth rates, (2) cell taxonomy, (3) ichthyotoxicity of monoclonal cultures at 25 and 35 psu in salinity, (4) differences in toxicity of lysed cells and supernatant at different cell concentrations (from 10 to  $100,000$  cells  $\text{ml}^{-1}$ ), and (5) temporal stability of lytic compounds. This study formally confirms the presence of *P. verruculosa* in Chilean waters using the large subunit (LSU) of the nuclear ribosomal RNA gene. The Chilean *P. verruculosa* ARC498 strain showed maximum cell densities at 30 psu (max.  $84,333 \pm 4,833$  cells  $\text{ml}^{-1}$ ) and maximum growth rates ( $\mu_{\text{max}}$ ) at 20 psu ( $1.44$  cells  $\text{d}^{-1}$ ). Cultures at 15 psu showed suppressed maximum cell density (max.  $269 \pm 71$  cells  $\text{ml}^{-1}$ ) but high  $\mu_{\text{max}}$  were recorded at the beginning of the exponential growth ( $1.31$  cells  $\text{d}^{-1}$ ). No significant differences were observed between lysed cells and supernatant treatments in the two salinity levels, suggesting that the most lytic portion is released into the cell-free media instead of remaining cell bound. Cytotoxicity was correlated to cell abundance, reducing gill cell viability down to 80 and 65% compared to controls at  $10,000$  and  $100,000$  cells  $\text{ml}^{-1}$ , respectively. Unexpectedly, lytic compounds from *P. verruculosa* ARC498 at 35 psu showed to be less toxic than cultures at 25, where a noticeable presence of peripheral mucocysts were observed by SEM. Lytic compounds from *in vitro* experiments are weakly toxic even at high cell concentrations, highly unstable and rapidly degraded in the light after 5 days of storage at  $15^{\circ}\text{C}$ . Our results point to the important effect of salinity on growth and ichthyotoxic potency of *Pseudochattonella* species and highlight the need for a deeper insight into the role of mucocysts in fish gill damage, which would provide a greater understanding as to the harmful modes of action of this species.

**Keywords:** molecular identification, fish-kills, fish cell line assay, ichthyotoxicity, Chilean fjords

## INTRODUCTION

Harmful Algal Blooms (HABs) have become an important threat for coastal aquaculture in the last decades (Hallegraeff et al., 2017). The more severe harmful effects occur when some microalgae produce toxic secondary metabolites that display potent biological activity against a wide variety of marine organisms or in humans upon consumption of contaminated seafood (Skjelbred et al., 2011). Among the ~80 species described as having the capacity to produce toxins, flagellates are recognized as the most harmful group (Wright and Cembella, 1998; Hallegraeff, 2003).

The heterokont genus *Pseudochattonella* encompass marine ichthyotoxic phytoflagellates recently added to the list of fish killing microalgae (Eckford-Soper and Daugbjerg, 2016b). Originally described as *Chattonella verruculosa* (Raphidophyceae) by Hara et al. (1994), molecular genetic analysis later on erected a new genus and *Pseudochattonella verruculosa* (Y. Hara and Chihara) Hosoi-Tanabe, Honda, Fukaya, Otake, Inagaki and Sako, was transferred to the class Dictyochophyceae (Hosoi-Tanabe et al., 2007). Subsequently a second species, *Pseudochattonella farcimen* (Eikrem, Edvardsen, and Throndsen), was assigned to this genus (Edvardsen et al., 2007; Eikrem et al., 2009). Blooms of *Pseudochattonella* species have caused several fish kills in Northern Europe (Lu and Goebel, 2000; Naustvoll et al., 2002; Edvardsen et al., 2007; Riisberg and Edvardsen, 2008), Japan (Yamamoto and Tanaka, 1990; Baba et al., 1995; Imai et al., 1998), New Zealand (MacKenzie et al., 2011; Chang et al., 2014) and Chile (Mardones et al., 2012; Clément et al., 2016; León-Muñoz et al., 2018).

The first reported bloom of *Pseudochattonella* sp. in Chilean waters occurred on January of 2004 in the locality of Cholgo – Los Lagos region (Mardones et al., 2012). *Pseudochattonella* sp. has since then been held responsible for several salmon mortality events in January and February of 2005, 2009, 2011, and in March 2016 (Mardones et al., 2012; Clément et al., 2016; León-Muñoz et al., 2018). The last bloom was the more massive one, extending several Km<sup>2</sup> in the inner zone of fjords and channels and causing the worst damage from HABs on Chilean salmon industry so far. Losses for the salmon industry in this occasion were calculated in 100,000 metric tons of Atlantic and Coho salmon and trout, equivalent to an export loss of US\$800 million (15% of Chile's yearly production) (Clément et al., 2016). This massive salmon mortality event evidenced the high fish-killing potency of the Chilean *Pseudochattonella* species, however, its ichthyotoxic mechanism is not yet understood.

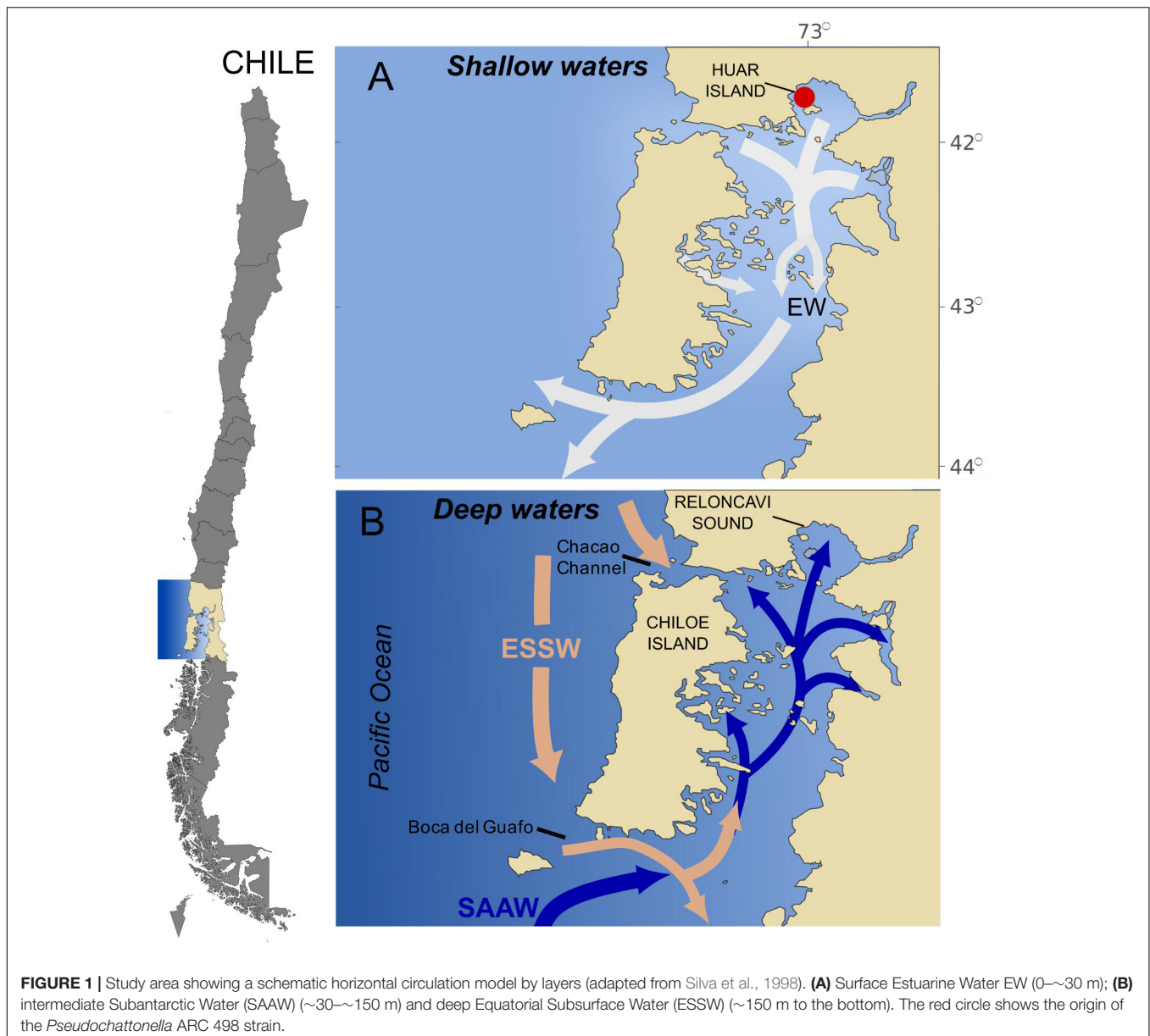
Research on *Pseudochattonella* spp. toxicity has proven to be difficult and has led to different opinions. Some studies have shown that cell culture extracts can produce a toxic effect on planktonic organisms and cell lines (Skjelbred et al., 2011; Chang et al., 2014), whereas studies conducted by Andersen et al. (2015) suggested that live cells are required to induce a toxic effect on fish. The latter hypothesis relates the toxicity of *Pseudochattonella* spp. to the presence of peripheral saccular extrusomes called “mucocysts” that

give these algae their characteristic warty appearance. The ecological function of the mucocysts is not clear but it has been suggested their implication as a mechanism to capture bacteria (Jeong et al., 2010) or as grazer deterrents (Tillmann and Reckermann, 2002).

The Chilean fjords are characterized by freshwater inputs from precipitation and melt water from Patagonian ice fields, which results in strong haline vertical gradients creating a unique highly productive area (Rignot et al., 2003). These stratified systems act as a barrier against the vertical propagation of turbulence from adjacent high mixing layers allowing for vertically heterogeneous phytoplankton distributions (Stacey et al., 2007; Yamazaki et al., 2010). Horizontally extensive subsurface patches of phytoplankton, the so-called “thin layers,” have been observed to persist from hours to weeks and can contain more than 75% of the total microalgae biomass in the water column (Holliday et al., 2003; Sullivan et al., 2010). Interestingly, key target HAB species in the northern Patagonian waters, such as *Dinophysis* spp. and the ichthyotoxic *Pseudochattonella* sp., have been observed in subsurface ‘thin layers’ (Alves-de-Souza et al., 2014; Clément et al., 2016). This fact points to the crucial role of salinity, as well as, nutrient availability and fluctuations in pCO<sub>2</sub>/pH for HAB formation in these highly heterogeneous estuarine systems.

The 2016 bloom seemed to be related to exceptional ocean and inshore water conditions in late 2015 due to a strong El Niño event and the positive phase of the Southern Annular Mode that altered the atmospheric circulation in the adjacent Pacific Ocean (León-Muñoz et al., 2018). The resultant increase in surface water temperature (>15°C) and reduced freshwater input allowed the advection of more saline (~33–35 in salinity) and nutrient-rich offshore waters into the fjords. These changes toward relatively high salinity in the Reloncaví Sound (25–27 psu) were observed prior to the bloom in February. By mid-March there was an increase in salinity (>30 psu) and a gradual drop in temperature, probably associated to lower river streamflow, which could have stimulated the massive *Pseudochattonella* bloom (León-Muñoz et al., 2018). The maximum cell concentrations during the event were estimated to be between 7,000 and 20,000 cells ml<sup>-1</sup> (Clément et al., 2016; León-Muñoz et al., 2018), several orders of magnitude higher of those recorded in previous *Pseudochattonella* sp. blooms in the area (Mardones et al., 2012). It has been suggested that high salinity variations in estuarine systems can strongly alter microalgae composition (Kirst, 1990), therefore this variable could have played key role in the proliferation, as well as, ichthyotoxic potency of *Pseudochattonella* sp. in the 2016 bloom event. However, ecophysiological studies on Chilean strains have not been yet undertaken since cultures of this species were only established after 2016 (Paredes et al., 2016; this study).

The aim of this study was to officially confirm the taxonomic identification of the Chilean *Pseudochattonella* populations, as well as, to provide the first assessment of growth parameters of species of this genus under different salinity levels. The potential lytic activity of *Pseudochattonella* species was also tested in intra- and extracellular metabolites using a fish gill cell line assay, as this has important implications for their mechanistic effects.



## MATERIALS AND METHODS

### Study Area

In the northern Patagonian fjords (42–44°S), where *Pseudochattonella* sp. bloom events have been registered, the vertical distribution of water masses presents a three-layer structure. A surface layer of Estuarine Waters (EW) (EW-Marine 21 to 31 psu; EW-Brackish 11 to 21 psu and EW-Freshwater 2 to 11 psu) between 0 and 20–30 m moves offshore increasing its salinity compared to fresh water sources (**Figure 1A**). An intermediate layer of Sub-Antarctic Waters (SAAW ~33 psu) between 30 and 150 m enters the inner part of the fjords from the south. The mix between EW and SAAW in the inner part of the fjords is called Modified Sub-Antarctic Waters (MSAAW). A deep layer of remnants of Equatorial Subsurface Waters

(ESSW > 33) from 150 m to the ocean bottom enters the inland sea of Chiloé through the Chacao channel and Boca del Guafo (**Figure 1B**; Silva et al., 1998). According to Castillo et al. (2016), the waters in the Reloncavi fjord are dominated along the seasons by EW in the upper layer and MSAAW in the deep layer.

### Culture Origin and Maintenance

#### Algal Culturing

A monoclonal culture of the toxic dictyochophyte *Pseudochattonella verruculosa* ARC498 strain was isolated from the Reloncavi sound during the 2016 bloom, kept in culture in the Algal Resources Collection at the University of North Carolina Wilmington and later transferred to the CREAN-IFOP laboratory to perform this study. Non-axenic cultures were grown in F/2 medium (Guillard and Ryther, 1962) at 15°C



in sterile filtered (0.22  $\mu\text{m}$ ) seawater (25 psu) at 20  $\mu\text{mol photon m}^{-2} \text{s}^{-1}$  (cool white fluorescent lamps) under a 12:12 h light:dark cycle.

### Gill Cell Line

The cell line RTgill-W1, originally cultured from gill filaments of rainbow trout *Oncorhynchus mykiss* (Bols et al., 1994), was acquired from the American Type Culture Collection (CRL-2523, ATCC). Cells were cultured at 17°C in the dark in Leibovitz's L-15 medium (L1518 Sigma), supplemented with 10% (v/v) fetal bovine serum (FBS, 12003C, Sigma), and an antibiotic-antimycotic solution (A5955, Sigma) containing amphotericin B (25  $\text{mg ml}^{-1}$ ), streptomycin (10  $\text{mg ml}^{-1}$ ) and penicillin (10,000 units  $\text{ml}^{-1}$ ) in 25- $\text{cm}^2$  culture-treated flasks (3100-025, Iwaki). 0.25% trypsin-0.02% EDTA in Hank's balanced salt solution (59428C, Sigma) was used to detach cells that grew as an adherent monolayer at the bottom of the flask. Subcultures were normally established twice per week at a ratio of 1:2 with L-15 medium renewal.

### Molecular Identification of *P. verruculosa*

Cells from monoclonal cultures of the ARC498 strain (GenBank accession number MK106355.1) were pelleted by centrifugation at 1500 g for 10 min at 4°C, the supernatant discarded and the cell pellets incubated at 65°C for 3 h in 1 ml CTB buffer and 10  $\mu\text{l}$  Proteinase K (10  $\text{mg/ml}$ ). The LSU (D1-D3) region of the rDNA gene was amplified using the primers D1R and D2C (Scholin et al., 1994; Edvardsen et al., 2003) in a SimpliAmp Thermo-Cycler (Applied Biosystems). The PCR protocol was as follow: initial denaturation step of 95°C for 1 min, followed by 35 cycles of 95°C for 30 s, 56°C for 30 s, 72°C for 30 s, followed by a 7 min extension at 72°C. PCR products were visualized on a 1.5% agarose gel and the product was purified using the Illustra GFX PCR DNA and gel band purification kit (GE Healthcare), and sequenced in both directions in Macrogen Korea. LSU rDNA sequences were aligned along with sequences available in GenBank using MUSCLE (Edgar, 2004), implemented in Geneious (Biomatters Ltd.). Unalienable regions were excluded from the LSU alignment. jModelTest (Posada, 2008) was used to identify appropriate models of sequence evolution (LSU: GTR + I), and maximum likelihood analyses were conducted using PhyML (Guindon et al., 2010). Support values were estimated using Likelihood Ratio Test (aLRT, Anisimova and Gascuel, 2006) and bootstrap analysis (1000 replicates). Bayesian analyses were conducted using MrBayes V3.1.2 (Ronquist and Huelsenbeck, 2003) using the appropriate model (GTR + I).

### Morphological Characteristics and Growth of *P. verruculosa* Cultures

#### Scanning Electron Microscopy (SEM) of *P. verruculosa*

Scanning electron microscopy analysis was performed following Botes et al. (2002). Briefly, a *P. verruculosa* culture was fixed 1:1 with 2% osmium tetroxide ( $\text{OsO}_4$ ) made up with filtered seawater. The 1% solution of fixed cells and  $\text{OsO}_4$  was applied on coverslips coated with poly-L-lysine (P-4832, Sigma) and left for ~30 min for *P. verruculosa* cells attachment. Attached cells were

washed by submerging the coverslips for 10 min in a 1:1 solution of distilled water and filtered seawater, followed for a 10-min wash in distilled water. The coverslips were then taken through a graded ethanol series (30, 50, 70, 80, 90, 95, and 100%, 10 min each step). Following removal from the 100% ethanol, few drops of hexamethyldisilazane (HMDS) were immediately added onto the coverslips to substitute critical point drying (CPD). Samples were coated with gold-palladium (60:40) and viewed with a JEOL JSM 6380LV SEM.

### *Pseudochattonella verruculosa* Growth

Exponentially growing cultures of *P. verruculosa* ARC498 were acclimated to the target salinity treatments of 15, 20, 25, 30, and 35 psu for 3 weeks prior to the experiments (corresponding to 10 generations). Pre-acclimated cultures were inoculated into sterile 60 ml polystyrene culture flasks containing 50 ml of F/2 media adjusted to the target salinity conditions. All treatments were performed in three replicates with an initial cell density of 500 cells  $\text{ml}^{-1}$ . The light intensity of 20  $\mu\text{mol photon m}^{-2} \text{s}^{-1}$  was chosen as it was considered the optimum saturation irradiance for the Chilean ARC498 *P. verruculosa* growth from pre-experimental measurements (data not shown). Thus, the growth rates obtained at the five salinity treatments correspond to maximum rates. The experiment was carried out for 27 days with samples taken every 3–7 days.

Cells concentration was determined immediately after sampling, based on buffered-Lugol's fixed cells quantified under an inverted light microscope (Olympus, CKX41) using a Sedgewick-Rafter chamber. In every sampling date, the mean cell number obtained from the three replicates was used to calculate the growth rate  $\mu$  ( $\text{d}^{-1}$ ):

$$\mu = \frac{\ln(c_1/c_0)}{t_1 - t_0} \quad (1)$$

where  $c_0$  and  $c_1$  are the cell concentrations (cell  $\text{ml}^{-1}$ ) at the beginning ( $t_0$ ) and end ( $t_1$ ) of the incubation period (days), respectively. At the end of the experiment, all cultures were left for 15 days under the original experimental conditions to observe the behavior of multinucleated cell aggregations as described by Tomas et al. (2004). These cell aggregations are believed to serve as resting stages and can potentially release a huge number of daughter cells that could alter the *in vitro* estimation of growth rates due to a sudden cell increase.

### *Pseudochattonella* Cytotoxicity Against Fish Gill Cells

The cytotoxicity assay with *P. verruculosa* ARC498 was carried out using conventional multiwell microplates according to Dorantes-Aranda et al. (2011). Cultures with confluent gill cells were trypsinized (59428C, Sigma) for detachment, counted using a hemocytometer and adjusted to a concentration of  $2 \times 10^5$  cell  $\text{ml}^{-1}$  in L-15 medium. Subsequently, gill cells were seeded in quadruplicate in 96-well flat bottom microplates (3860-096, Iwaki, Japan), using a volume of 100  $\mu\text{l}$  per well. After 48 h at 17°C in the dark for gill cell attachment, L-15 medium was discarded, the cells rinsed with PBS and exposed to 100  $\mu\text{l}$

*Pseudochattonella* lysed cells and supernatant medium from cultures at 25 and 35 psu at 17°C in the dark. A salinity of 25 corresponds to a value measured above the pycnocline during the 2016 El Niño event in the Reloncaví Sound (Clément et al., 2016), and 35 psu is consistent with salinity values measured in the offshore Equatorial Subsurface Water (ESSW) (Silva et al., 1998). Supernatant and lysed cells treatments were prepared from *Pseudochattonella* cultures in the exponential growth phase at five different concentrations (10, 100, 1000, 10,000, and 100,000 cells ml<sup>-1</sup>). The supernatant suspension was prepared by centrifugation of cultures for 5 min at 3500 rpm and then diluted as needed. The lysed cell suspension was prepared by sonication of diluted samples for 10 min at an amplitude of 10 µm peak to peak at 17°C and filtered using a syringe with a nylon filter (0.22 µm). After 1 h exposure, the viability of the gill cells was determined using L-15/ex medium (Schirmer et al., 1997), a modified version of the L-15 medium, containing 5% of the indicator dye alamarBlue (DAL1025, Invitrogen) (Pagé et al., 1993). The medium containing the indicator dye was added to all cell-seeded wells and incubated for 2 h in the dark (Dayeh et al., 2005). Using a microplate reader (FLUOstar OMEGA, BMG Labtech), the fluorescence signal of alamarBlue was detected using excitation and emission filters of 540 and 570 nm, respectively. The viability of the gill cells was expressed as response percentage of the treatments relative to the controls (% of control).

## Data Analysis

To explore the effect of salinity on growth and the ichthyotoxic potency of *P. verruculosa*, analysis of variance (ANOVA) from simple linear regression models on maximum cell density, maximum specific growth rate ( $\mu_{\max}$ ) and gill cell viability against *Pseudochattonella* cell density and intra- and extracellular compounds was performed. Normality and homogeneity of variances were assessed by the Kolmogorov-Smirnov method and Levene's test. A *post hoc* analysis using a Tukey HSD test was performed to determine differences among treatments. The null hypothesis (no difference in responses) was rejected in all statistical analyses if the respective *p*-value was <0.05. These analyses were performed using the software R v. 3.0.1 (Ihaka and Gentleman, 1996<sup>1</sup>).

## RESULTS

### Molecular Identification of *Pseudochattonella*

The 250 bp partial LSU rDNA sequence obtained from *P. verruculosa* ARC498 was identical to six sequences attributed to *P. verruculosa* (AB217643.1, AM850226.1, AM850225.1, AM850224.1, AB217642.1, and AM040504.1), and differed from sequences of *Pseudochattonella farcimen* by 4–5 substitutions. Phylogenetic trees obtained from these sequences and outgroup rooted with relatives of *Pseudochattonella* species demonstrate this pattern and give a relative scale of variation between the

Chilean strain and other *P. verruculosa* sequences as well as *P. farcimen*. The LSU tree shows the Isla Huar isolate resolved in the *P. verruculosa* clade, although not as a monophyletic group (Figure 2).

### Morphological Characteristics and Growth of *P. verruculosa* Cultures

Cells showed a variable morphology with form and size changing in response to growth phase and salinity treatments. All treatments showed cell shapes ranging from small spherical/oval (~4–10 µm) to conical-elongated (~8–19 µm). Two heterokont flagella were present. The shortest flagellum faces backward and a longer anteriorly directed flagellum (~12–22 µm) pulls the cell forward (Figure 3A). Mucocysts were mainly present in larger than smaller cells. SEM analysis micrographs of the *P. verruculosa* ARC498 cultures (used for the gill cell line assay) showed a more recurrent presence of mucocysts in *P. verruculosa* cells at 25 psu than cells at 35 psu (Figures 3A,B). Mucus secretion was observed to form cell aggregations (Figure 3C).

The response of *P. verruculosa* ARC498 to changes in salinity was significantly different in terms of maximum cell density and maximum specific growth rate ( $\mu_{\max}$ ) (ANOVA, *df* = 4, *p* < 0.001, Figure 4). The highest maximum cell density among all treatments was obtained at salinity 30 (84333 ± 4833 cells ml<sup>-1</sup>), and the lowest at salinity 15 (269 ± 71 cells ml<sup>-1</sup>) (Table 1). The highest  $\mu_{\max}$  of 1.44 days<sup>-1</sup> was reached at 20 psu and the lowest at 30 psu with a mean value of 0.88 days<sup>-1</sup>. Multinucleated cell aggregations were observed at the bottom of the flasks throughout the experimental period. After ~15 days, some cultures exhibited vegetative cell growth that reached modest concentrations under depleted-nutrient conditions (data not shown).

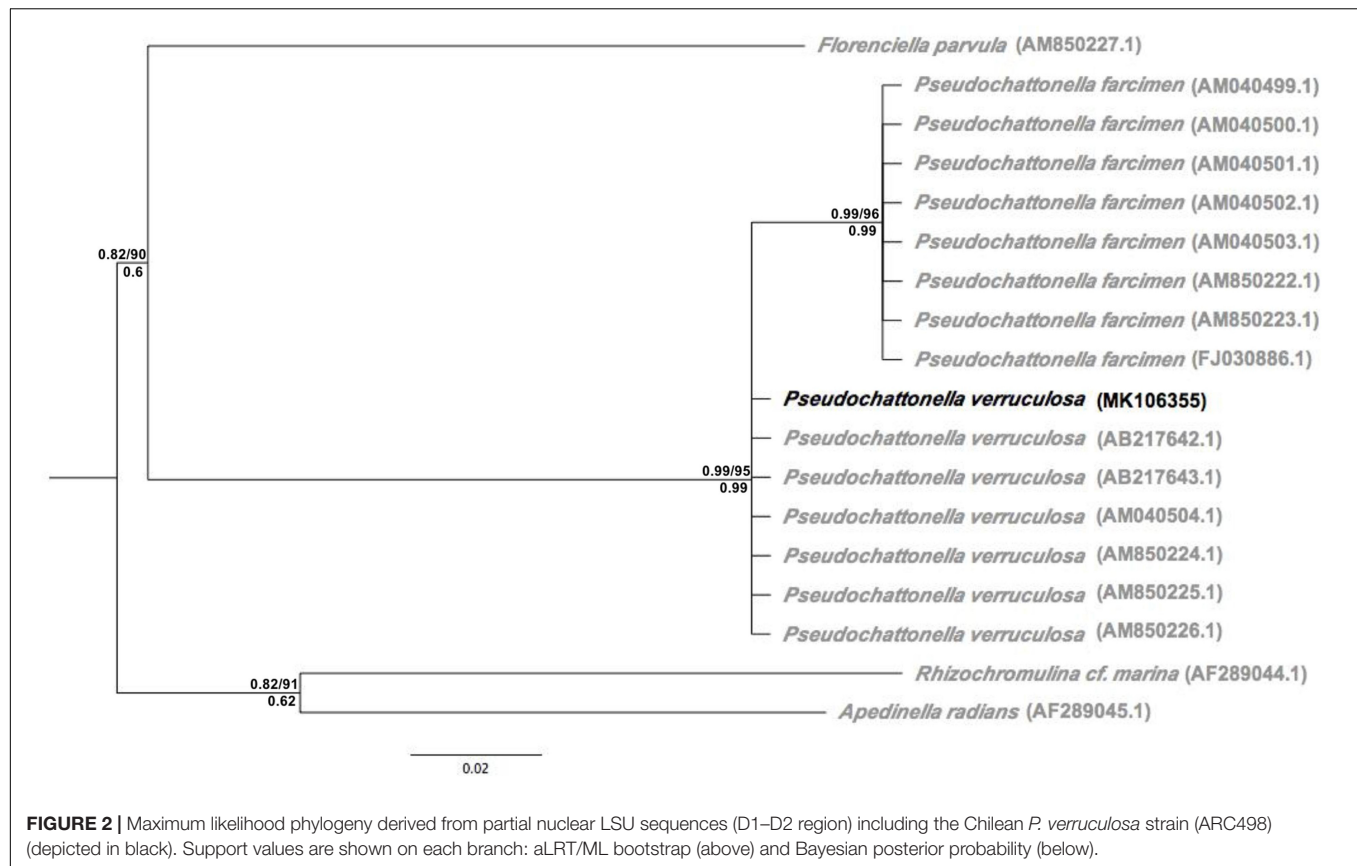
### *P. verruculosa* Cytotoxicity Against Fish Gill Cells

Viability of gill cells after 1 h exposure to *P. verruculosa* did not show significant differences between the supernatant and ruptured cell treatments (ANOVA, *p* > 0.05, Figure 5), and lytic activity showed to slightly increase together with *Pseudochattonella* cell density at both salinity treatments. Cytotoxicity of intra- and extracellular compounds of *P. verruculosa* did not show significant differences (ANOVA, *p* < 0.05; except treatment at 100,000 *Pseudochattonella* cells ml<sup>-1</sup> at 25 psu-day1; Figure 5A) and treatments at salinity 25 showed to be more potent than toxic compounds obtained at salinity 35, reducing gill cell viability down to 52% of controls at 100,000 cells ml<sup>-1</sup>. Lytic compounds of both salinity treatments rapidly degraded in the light conditions, reducing gill cell viability to only less than 20% after 5-days storage at 15°C (Figure 5).

## DISCUSSION

As *P. verruculosa* is a newly reported species for Chilean waters, almost nothing is known about its ecology. This is the first study to formally confirm the taxonomic identification of Chilean

<sup>1</sup><http://www.r-project.org>



*Pseudochattonella* populations as *P. verruculosa* using a genetic approach and the first assessment of *in vitro* cell growth and cytotoxicity.

## Molecular Characterization

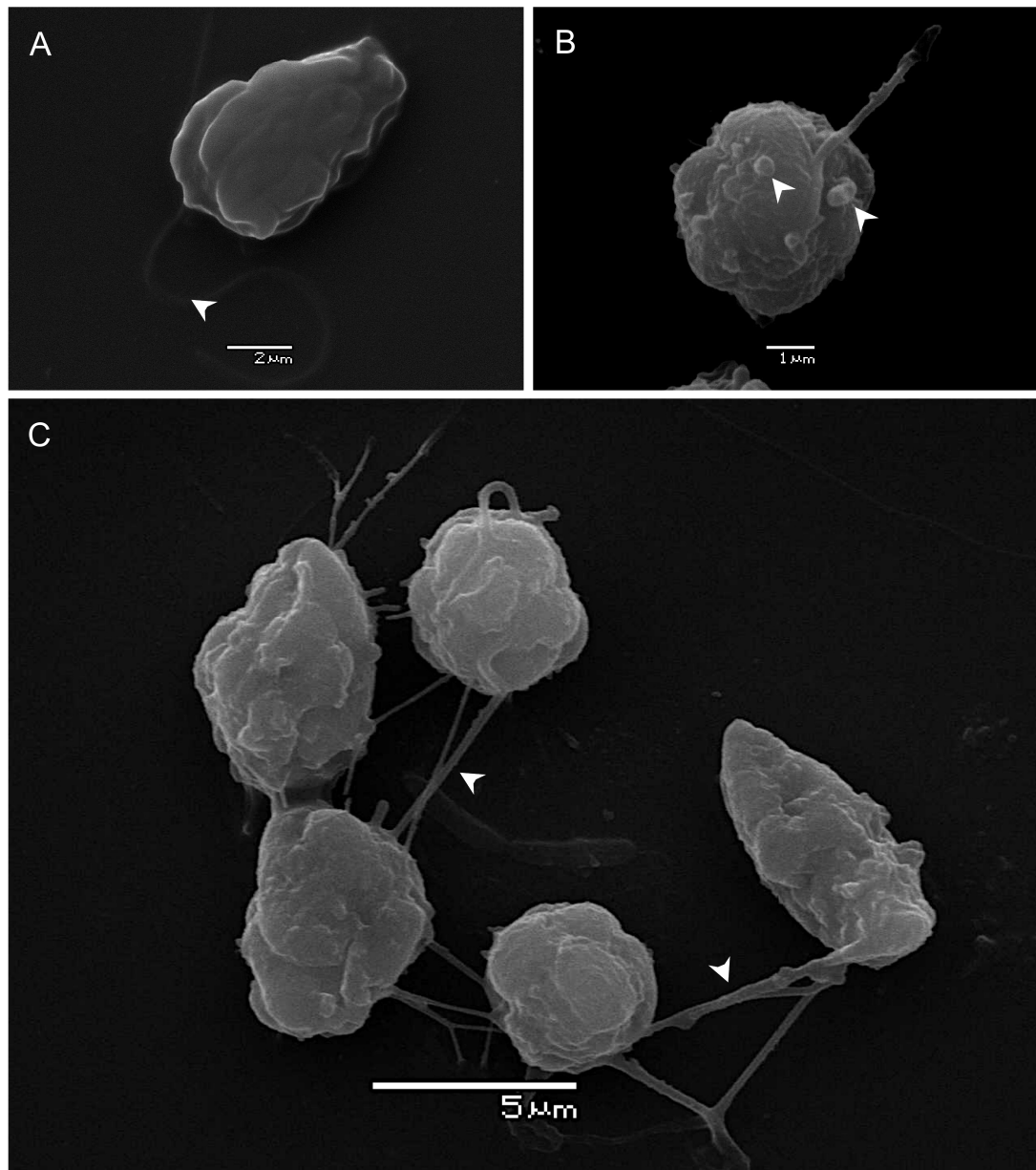
To the best of our knowledge, this is the first formal molecular characterization of a *Pseudochattonella* strain isolated from the massive 2016 bloom in the Reloncaví sound (southern Chile), using a sequence obtained from the LSU rDNA region (Figure 1). These results resolved the Chilean *P. verruculosa* strain (ARC498) within a clade with others *P. verruculosa* strains from Japan and New Zealand in agreement with Chang et al. (2014) for the same gene. However, other markers (18S, RbcL, COI, and ITS) should be used to confirm our Chilean *Pseudochattonella* strain within the *P. verruculosa* clade. This is mainly due to the fact that different gene markers can differ in results. For instance, a *Pseudochattonella* strain Wellington Harbour (New Zealand) was found to be similar to *P. verruculosa* using the large subunit of ribulose biphosphate carboxylase (rbcL) and the partial sequences of the nuclear encoded LSU rDNA, whereas sequences of the mitochondrial cytochrome c oxidase subunit (COI) gene grouped this strain as *P. farcimen* (Chang et al., 2014). Moreover, unpublished sequence data from field samples collected during the 2016 bloom have indicated the potential co-existence of *P. farcimen* blooming together with *P. verruculosa* in the Patagonian fjord area (ADL Diagnostic Chile- personal communication). Under this scenario and given the high

morphological variability already described for *Pseudochattonella* species (Eckford-Soper and Daugbjerg, 2016b), the description of molecular markers becomes a key tool for precise species detection in the Chilean waters. The *P. verruculosa* ARC498 sequence described in this study has the potential for the design of primers for quantitative PCR (qPCR) aiming a rapid and low-cost identification and quantification tool as used in field samples from *Pseudochattonella* sp. blooms in Danish waters (Eckford-Soper and Daugbjerg, 2016a). In future studies, HPLC pigment analysis could also help for a better *Pseudochattonella* species discrimination since violaxanthin, lutein, and antheroxanthin are only detected in *P. verruculosa* (Tomas et al., 2004).

## Effect of Salinity on *P. verruculosa* Cultures

High salinity variations observed in estuarine systems due to water mix or stratification can produce alterations in marine microalgae composition. These effects can include (i) changes in the cellular ionic ratios due to the selective ion permeability of the membrane, (ii) osmotic stress with direct impact on the cellular water potential and (iii) ionic stress caused by the unavoidable uptake or loss of ions (Kirst, 1990). The Reloncaví Sound (41°35'S, 72°20'W), which is recognized as the 'hot spot' for *Pseudochattonella* outbreaks in the south of Chile, has a broad range of salinity that can vary from <10 psu at the surface to >30 psu in deep waters (Castillo et al., 2016). In our study, the *P. verruculosa* ARC498 strain



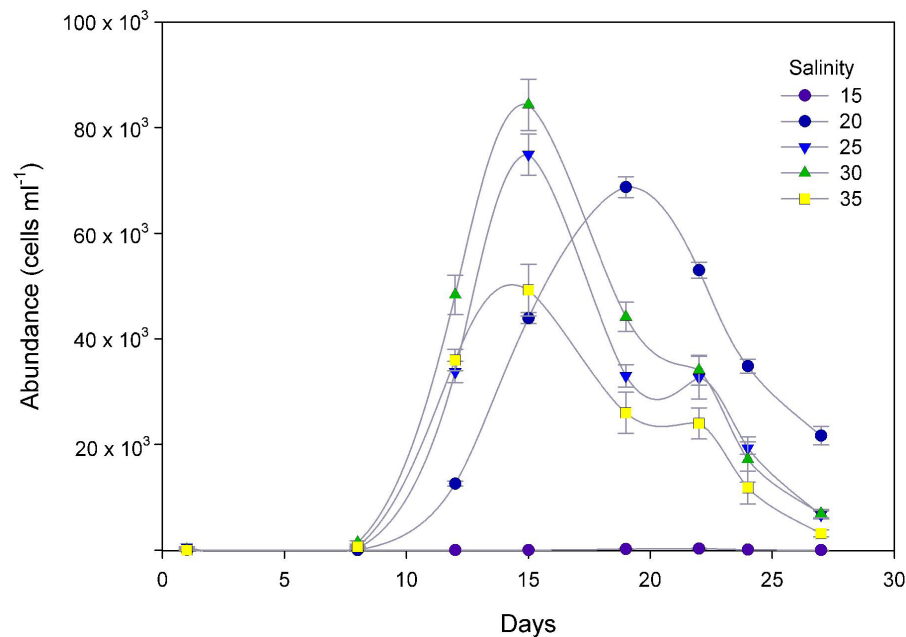


**FIGURE 3 |** Scanning electron microscopy of *P. verruculosa* ARC498 strain. **(A)** Ovoid cell with a long forwardly directed flagellum (arrowhead) from culture at 35 in salinity, **(B)** spherical cell with mucocysts at the periphery of the cell (arrowheads) from culture at 25 in salinity, and **(C)** group of cells interconnected by thin mucus-filaments (arrowheads).

isolated from the Reloncaví sound reached high cell densities between 20 and 35 psu. At salinity 15, exponential cell growth was 8 days delayed compared to the other salinity treatments and cell densities were several orders of magnitude lower, although positive net growth lasted for several days. These results suggest that freshwater inputs in the upper layer of the fjord may act as an environmental control for *P. verruculosa* cell growth.

The capacity to maintain high and positive net growth at a broad range of salinity values may have several competitive

advantages in a dynamic estuarine system when turbulence breaks the halocline and vertically disperse cell through the water column. This pattern of broad salinity optima has also been observed in the distantly related *Dictyocha speculum* Ehrenberg (Henriksen et al., 1993). However, it has been suggested that microalgal cell growth primarily depends on temperature and light, with the tolerated salinity range becoming broader as these parameters approach optimality (Kirst, 1990). An increasing tolerance to a high salinity range by our *P. verruculosa* strain could be then a result of *in vitro* optimal conditions (20 μmol



**FIGURE 4 |** Temporal cell growth curves of *P. verruculosa* cultures under five salinity treatments. Symbols represent the mean and error bars the standard deviation of cell counts from triplicate measurements.

photon  $\text{m}^{-2} \text{s}^{-1}$  and  $15^{\circ}\text{C}$ ) and it is likely to change in the highly dynamic fjords. Other aspect to take into account is that the experiments were performed using strains previously acclimated for several generations. Further studies should assess the effect of drastic salinity changes on *P. verruculosa* growth.

In this study, we documented for the first time the cell growth rates for a Chilean *P. verruculosa* strain. Maximum growth rates that ranged between  $0.88$  to  $1.44 \text{ days}^{-1}$  were higher compared to those observed by Skjelbred et al. (2013) for Danish *P. verruculosa* strains ( $0.61 \text{ days}^{-1}$ ) and lower than  $1.74 \text{ days}^{-1}$ , which was recorded for a Japanese strain (Yamaguchi et al., 1997). The relative high  $\mu_{\text{max}}$  observed in our experiment could be attributed to a high cell division rate but also to the presence of multinucleated cell aggregations noted at the bottom of our treatment flasks throughout and  $\sim 15$  days after the end of the experiment. Similar *in vitro* cell aggregations have also been observed in other *P. verruculosa* strains from the North Sea and Japan (Tomas et al., 2004). It has been suggested that these massive aggregations could act as resting stages when conditions become unfavorable (Jakobsen et al., 2012; Chang et al., 2014). Thus, *in vitro* growth rate values could be enhanced by germling cell input from resting stages and also might point to homothallism in our monoclonal ARC498 strain.

### Lytic Activity of *P. verruculosa*

Species of the genus *Pseudochattonella* has been reported to be highly ichthyotoxic with important economic losses for the Chilean salmon industry since 2004 (Mardones et al., 2012). Most examinations of affected salmon during natural fish-kill events in Chile have shown noticeable edema, hyperplasia and

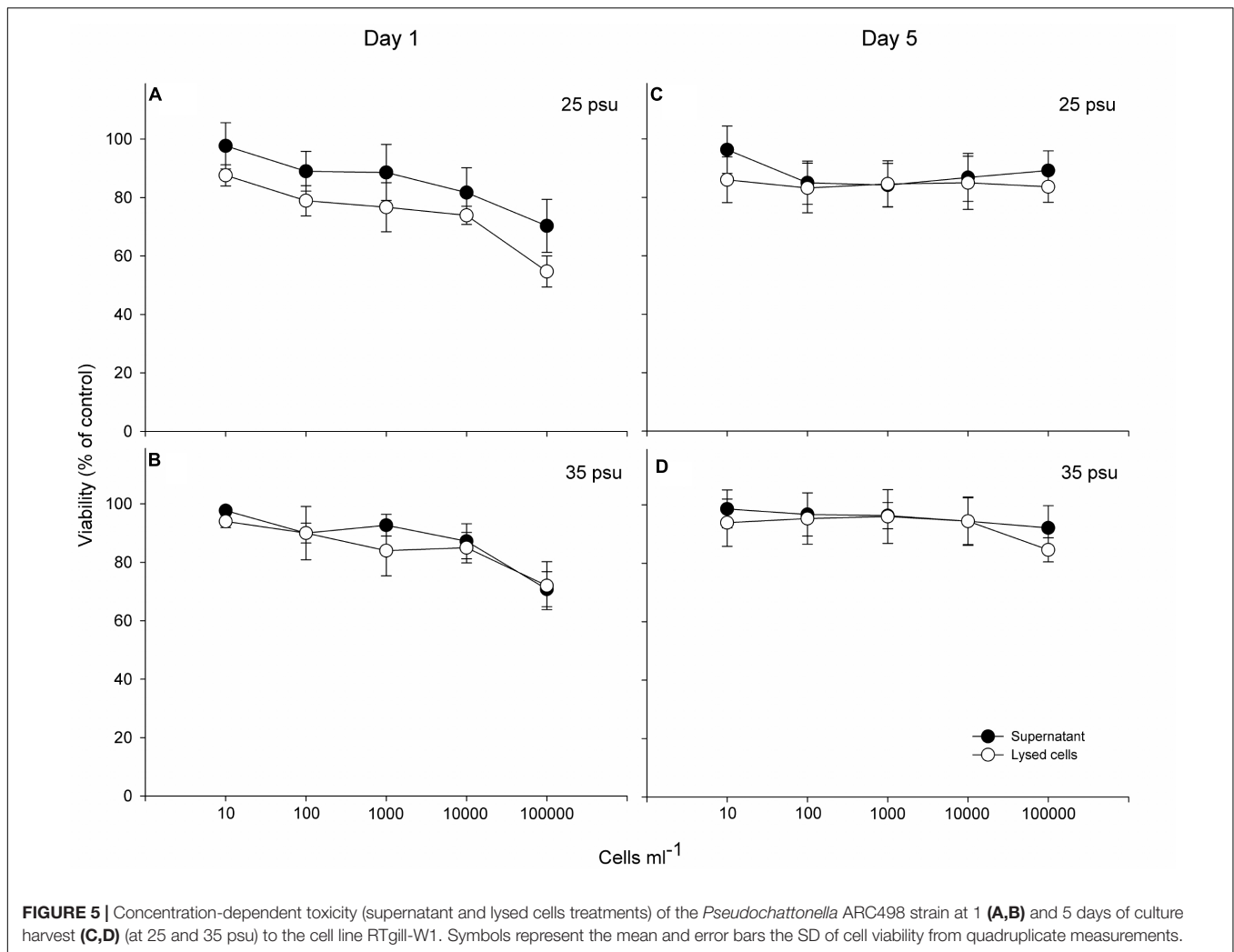
**TABLE 1 |** Growth parameters of *Pseudochattonella verruculosa* cultures (mean  $\pm$  SD) maximum cell density and  $\mu_{\text{max}}$  mean maximum specific growth rate and under five different salinity treatments.

Treatment (salinity)	Maximum cell density (cells $\text{ml}^{-1}$ )	$\mu_{\text{max}}$ ( $\text{d}^{-1}$ )
15	$269 \pm 71$	1.31
20	$68750 \pm 1963$	1.44
25	$74916 \pm 3892$	1.12
30	$84333 \pm 4833$	0.88
35	$49250 \pm 4793$	1.01

necrosis of secondary gill lamellae (C. Sandoval – VEHICE, personal communication). Despite this, research on the toxic compounds produced by this species is incipient and has proven to be difficult.

The fish cell line RTgill-W1 assay has been widely used to assess cell viability in ichthyotoxic-microalgal studies (Dorantes-Aranda et al., 2015). In this work, we experimentally demonstrated lytic activity of the Chilean *P. verruculosa* ARC498 strain toward the RTgill-W1 cells, although ecologically realistic *P. verruculosa* bloom concentrations ( $10,000$ – $100,000$  cells  $\text{ml}^{-1}$ ) exhibited only very limited loss of viability (down to max. 45% of controls). These results differ from field observations in the Chilean bloom, where *P. cf. verruculosa* showed to be extremely toxic at cell concentrations as low as  $5$  cells  $\text{ml}^{-1}$  (Mardones et al., 2012). Similarly, Montes et al. (2018), based on Chilean phytoplankton data bases, estimated that  $<1$  cell  $\text{ml}^{-1}$  can be associated with anomalous salmon behavior in the Patagonian fjords. Comparable to these observations in the Chilean waters, only  $10$  cells  $\text{ml}^{-1}$  of *P. verruculosa*





were enough to cause mortality of salmon in New Zealand (MacKenzie et al., 2011). There might be a few experimental conditions that can explain our findings: (1) gill cells were shortly exposed to *P. verruculosa* toxic compounds (1 h) compared to other studies (Dorantes-Aranda et al., 2015; Mardones et al., 2015). Mardones et al. (2015) showed that some strains of the toxic dinoflagellate *Alexandrium catenella* were able to reduce gill cell viability down to 20% of controls at 4,000 cells ml<sup>-1</sup> after 2 h exposure due to net K<sup>+</sup> efflux from fish gill cells (Mardones et al., 2018). Thus, an increment in the time of exposure to *P. verruculosa* toxic compounds is likely to produce a more severe gill tissue damage; (2) in this study lytic compounds from *P. verruculosa* treatments were tested against the gill cells at 1 and 5 days after their extraction. This storage procedure showed that cytotoxic potency rapidly vanished over a 5-day period. It is likely therefore that *P. verruculosa* exudates could be more ichthyotoxic as soon as they are released into the surrounding marine environment. This early toxic effect was previously observed in *in vitro* cytotoxic assays using algal extracts of *Pseudochattonella* species (Skjelbred et al., 2011); (3) live cells are required to induce

a toxic effect on fish as suggested by Andersen et al. (2015); and/or (4) the reduced cytotoxicity observed on gill cells could be merely due to the fact that the ARC498 strain is not highly toxic.

The ichthyotoxic mechanism in *Pseudochattonella* species is elusive with the mode of action yet to be completely determined (Eckford-Soper and Daugbjerg, 2016b). In this study, cytotoxicity of lysed *P. verruculosa* cells did not show significant differences to that of the supernatant. This suggests that the most lytic portion is released into the cell-free media instead of remaining cell bound. These results might lead to a future explanation of the nature of the toxins produced by *Pseudochattonella* species and their role in chemical ecology. Initial ideas focused on the concept that phycotoxins acted as defensive chemicals against competitors and predators (Smetacek, 2001). Modern studies suggest that cell bound phycotoxins might play a different ecological function compared to secondary toxic metabolites that are released from the cell to the surrounding environment. Excretion of a toxic compound by individual cells may aim to cease grazing by a motile predator or produce an antibiotic effect to reduce the colonization of microalgal cells by viruses

or bacteria (Cembella, 2003). For instance, allelochemical activity by the dinoflagellate *Alexandrium tamarense* against the predatory dinoflagellate *Oxyrrhis marina* was shown to be unrelated to paralytic shellfish toxins (PST) (Alpermann et al., 2010). Similarly, Mardones et al. (2015, 2018) showed that PST are not involved in fish gill damage, but rather an alternative ichthyotoxic mechanism was suggested based on lipid peroxidation products due to a synergistic interaction between Reactive Oxygen Species (ROS) and Polyunsaturated Fatty Acids (PUFA). Interestingly, *Pseudochattonella* species have shown high proportions of a rare PUFA C18:5n-3 and C16:1n5 and high ratios of docosahexaenoic acid (DHA) to eicosapentaenoic acid (EPA) (Giner et al., 2008; Dittami and Edvardsen, 2012). This rises a big question about the potential involvement of a PUFA/ROS synergism in the potent ichthyotoxicity observed for the Chilean *P. verruculosa* ARC498. Key research on PUFAs cell content, as well as, ROS and other active secondary metabolites production for Chilean strains remain to be conducted.

Salinity had a significant effect on *P. verruculosa* ichthyotoxic potency. Cultures at salinity 25 were more toxic than cultures at 35 psu. It was also observed that *Pseudochattonella* cells at salinity 25 were bigger and showed a higher presence of mucocysts than cells grown at salinity 35. These mucocysts are present in several species of raphidophytes that have been related to fish-killing events, e.g., *Heterosigma akashiwo* (Twiner et al., 2005) and *Fibrocapsa japonica* (Pezzolesi et al., 2010). This suggests the putative involvement of mucocysts to induce a toxic effect on fish gill tissue as suggested by Andersen et al. (2015). However, no experiments to test cytotoxicity with whole cells of Chilean *Pseudochattonella* have been performed yet.

## Ecological Implications

In this study, changes in salinity were an important driver for *in vitro* *Pseudochattonella* cell growth and ichthyotoxic potency. The enhanced cell growth under moderate-high salinity conditions ( $\sim 30$ ) is in line with field observations during the massive 2016 *Pseudochattonella* sp. outbreak. The strong El Niño 2015–2016 resulted in an extremely dry summer in the southeast Pacific coast with a record low streamflow and higher than normal solar radiation (León-Muñoz et al., 2018). These extreme meteorological conditions allowed vertical advection of saline and nutrient-rich waters ( $\sim 30$  psu) that ultimately resulted in an enhanced *Pseudochattonella* sp. bloom that reached peak cell densities of  $\sim 7,700$  to  $20,000$  cells  $\text{ml}^{-1}$  (Clément et al., 2016; León-Muñoz et al., 2018). Thus, the lower *Pseudochattonella* cell concentrations recorded in the 2005, 2009, and 2011 outbreaks could have been the result of highly stratified water columns with higher freshwater inputs in the surface. However, those low-cell density outbreaks produced substantial fish gill damage with less than  $20$  cells  $\text{ml}^{-1}$ , suggesting the important effect of moderate-lower salinity on the enhanced ichthyotoxic potency ( $\sim 20$ – $25$  psu).

Although the macro-scale El Niño 2015–2016 event produced evident changes in the haline structure of the water column in the Reloncaví Sound, it is important to highlight that  $p\text{CO}_2/\text{pH}$  and nutrient composition/availability, among other

variables, should also have been affected. Mardones et al. (2016) showed that spatio-temporal  $p\text{CO}_2/\text{pH}$  fluctuations in Chilean fjords can strongly affect physiological responses in *A. catenella*. Physiological changes such as reduced cell size and enhanced chain formation at high pH/low  $p\text{CO}_2$  (i.e., phytoplankton bloom conditions) were observed. On the other hand, the entry of offshore waters rich in inorganic nutrients (i.e.,  $\text{NO}_3^-$ ) vs. inland waters, highly influenced by organic nutrients (i.e.,  $\text{NH}_4^+$ ) due to intense aquaculture, might have contributed to changes in phytoplankton community composition by differences in  $\text{NH}_4^+$  and  $\text{NO}_3^-$  taxon-specific metabolism. The role of these key chemical variables in the bloom dynamics of *Pseudochattonella* species should be pursued in future experiments.

Finally, the extremely high cell densities obtained in our *in vitro* experiments ( $>80,000$  cells  $\text{ml}^{-1}$ ) compared to field counts, may reflect the difficulty in the *Pseudochattonella* sp. cell identification using light microscopy. The complex life cycle of *Pseudochattonella* species, that also includes very small flagellate and large multinucleated stages (Chang et al., 2014; this study), likely lead to misidentification and cell count sub-estimations of the 2016 *Pseudochattonella* sp. outbreak. Therefore, the advent of molecular approaches becomes an important tool for improving the effectiveness of monitoring programs.

In conclusion, the present work has formally confirmed the presence of *P. verruculosa* in Chilean waters, as well as, showed the effect of salinity on cell growth and ichthyotoxic potency. Overall, high growth rates observed at moderate-high salinity might help to explain the massive outbreak recorded in 2016. The increasing cytotoxicity by *P. verruculosa* under low-salinity conditions may be important to understand fish-killing properties of the Chilean strains since blooms of this dictyochophyte occur in highly variable oceanographic conditions, such as those occurring in the southern Chilean coast. The role of mucocysts, as well as, production of PUFAs, ROS and/or secondary metabolites that could explain ichthyotoxicity by Chilean strains of *P. verruculosa* remain to be investigated.

## AUTHOR CONTRIBUTIONS

All authors conceived, designed, and performed the experiments and analysis. JM analyzed the data and wrote the manuscript with contributions from all authors.

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**Conflict of Interest Statement:** The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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